



## PATENT ABSTRACTS OF JAPAN

(11) Publication number: **05117268 A**(43) Date of publication of application: **14.05.93**

(51) Int. Cl.

**C07D401/12****A61K 31/44****A61K 31/47****C07D401/14**

**/(C07D401/12 , C07D213:00 ,  
C07D235:00 ), (C07D401/14 , C07D213:00  
, C07D217:00 , C07D235:00 )**

(21) Application number: **03304219**(22) Date of filing: **22.10.91**(71) Applicant: **YOSHITOMI PHARMACEUT IND LTD**

(72) Inventor: **KAWAKITA TAKESHI  
YAMAGUCHI HIROKO  
HAGA KEIICHIRO  
IKEDA TAKASHI  
YOKOYAMA YOSHIHITO**

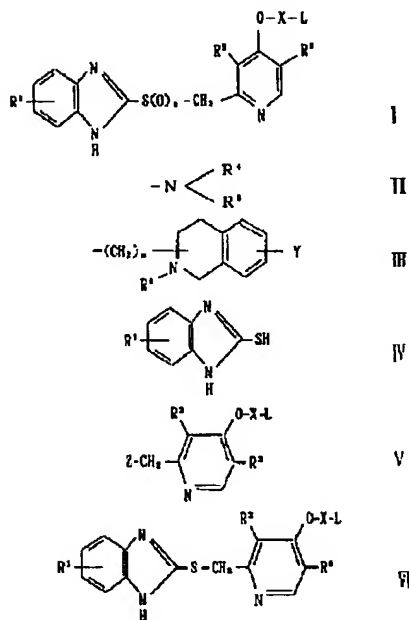
(54) **PYRIDINE COMPOUND**

(57) Abstract:

PURPOSE: To obtain a new compound useful as a during for antiulcer action, gastrointestinal cell protecting action, action for preventing recrudescence and palindromia of ulcer, antimicrobial action against *Helicobacter pylori*.

CONSTITUTION: A compound of formula I [ $R^1$  is H, halogen, alkyl, alkoxy, alkoxycarbonyl or haloalkyl; (n) is 0-2;  $R^2$  and  $R^3$  are H, halogen or alkyl; X is alkylene; L is formula II ( $R^4$  and  $R^5$  are alkyl, cycloalkyl, etc.; X-L is formula III [(m) is 1-3;  $R^6$  is phenylalkyl; Y is H, halogen, alkyl, etc.], e.g. 2-[3-methyl-4-(2-(N-benzyl-N-cyclohexylamino)ethoxy)pyridyl]-methylthio-1H-benzimidazole. The compound is obtained by reacting a compound of formula IV with a compound of formula V (Z is halogen, sulfonyloxy, etc.) and oxidizing the resultant compound of formula VI.

COPYRIGHT: (C)1993,JPO&amp;Japio


**BEST AVAILABLE COPY**